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Neue für die Gene metR und metZ kodierende Nukleotidsequenzen

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By: \_\_\_\_\_

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## Certificate of Priority for Filing of a Patent Application

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**Title:** New nucleotide sequences which code  
for the metR and metZ genes

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Faust

**New nucleotide sequences which code for the metR  
and metZ genes**

The invention provides nucleotide sequences from coryneform bacteria which code for the metR and metZ genes and a  
5 process for the fermentative preparation of amino acids, in particular L-methionine, by attenuation of the metR and/or metZ gene.

Prior art

10 L-Amino acids, in particular methionine, are used in human medicine and in the pharmaceuticals industry, in the foodstuffs industry and very particularly in animal nutrition.

It is known that amino acids are prepared by fermentation from strains of coryneform bacteria, in particular  
15 Corynebacterium glutamicum. Because of their great importance, work is constantly being undertaken to improve the preparation processes. Improvements to the process can relate to fermentation measures, such as, for example, stirring and supply of oxygen, or the composition of the  
20 nutrient media, such as, for example, the sugar concentration during the fermentation, or the working up to the product form by, for example, ion exchange chromatography, or the intrinsic output properties of the microorganism itself.

25 Methods of mutagenesis, selection and mutant selection are used to improve the output properties of these microorganisms. Strains which are resistant to antimetabolites or are auxotrophic for metabolites of regulatory importance and which produce amino acids, such  
30 as e.g. L-methionine, are obtained in this manner.

Methods of the recombinant DNA technique have also been employed for some years for improving the strain of Corynebacterium strains which produce L-amino acids, by

amplifying individual amino acid biosynthesis genes and investigating the effect on the amino acid production.

#### Object of the invention

The inventors had the object of providing new measures for improved fermentative preparation of amino acids, in particular L-methionine.

#### Description of the invention

Where L-amino acids or amino acids are mentioned in the following, this means one or more amino acids, including their salts, chosen from the group consisting of L-asparagine, L-threonine, L-serine, L-glutamate, L-glycine, L-alanine, L-cysteine, L-valine, L-methionine, L-isoleucine, L-leucine, L-tyrosine, L-phenylalanine, L-histidine, L-lysine, L-tryptophan and L-arginine.

When L-methionine or methionine are mentioned in the following, the salts, such as e.g. methionine hydrochloride or methionine sulfate are also meant by this.

The invention provides isolated polynucleotides from coryneform bacteria, which comprise the polynucleotide sequences which code for the metR and/or metZ genes, chosen from the group consisting of

- a) polynucleotide which is identical to the extent of at least 70% to a polynucleotide which codes for a polypeptide which comprises the amino acid sequence of SEQ ID No. 2,
- b) polynucleotide which is identical to the extent of at least 70% to a polynucleotide which codes for a polypeptide which comprises the amino acid sequence of SEQ ID No. 3,
- c) polynucleotide which codes for a polypeptide which comprises an amino acid sequence which is identical to

the extent of at least 70% to the amino acid sequence of  
SEQ ID No. 2,

- d) polynucleotide which codes for a polypeptide which  
comprises an amino acid sequence which is identical to  
5 the extent of at least 70% to the amino acid sequence of  
SEQ ID No. 3,
- e) polynucleotide which is complementary to the  
polynucleotides of a), b), c) or d), and
- f) polynucleotide comprising at least 15 successive  
10 nucleotides of the polynucleotide sequence of a), b),  
c), d) or e),

the polypeptides according to a) or c) preferably having  
the activity of the transcription activator MetR and the  
polypeptides according to b) or d) preferably having the  
15 activity of O-succinylhomoserine sulfhydrylase (MetZ).

The invention also provides the abovementioned  
polynucleotides, these preferably being DNAs which are  
capable of replication, comprising:

- (i) the nucleotide sequence shown in SEQ ID No. 1, or
- 20 (ii) at least one sequence which corresponds to  
sequence (i) within the range of the degeneration  
of the genetic code, or
- (iii) at least one sequence which hybridizes with the  
sequences complementary to sequences (i) or (ii),  
25 and optionally
- (iv) sense mutations of neutral function in (i).

The invention also provides:

a DNA which is capable of replication and comprises the  
nucleotide sequence as shown in SEQ ID No. 1,

a polynucleotide which codes for a polypeptide which comprises the amino acid sequence as shown in SEQ ID No. 2 or SEQ ID No. 3,

5 a vector containing parts of the polynucleotide according to the invention, but at least 15 successive nucleotides of the sequence claimed

and coryneform bacteria in which the metR gene and/or the metZ gene is or are attenuated, in particular by deletion, insertion or base exchange.

10 The invention also provides polynucleotides which substantially comprise a polynucleotide sequence, which are obtainable by screening by means of hybridization of a corresponding gene library of a coryneform bacterium, which comprises the complete gene or parts thereof, with a probe  
15 which comprises the sequence of the polynucleotide according to the invention according to SEQ ID No. 1 or a fragment thereof, and isolation of the polynucleotide sequence mentioned.

Polynucleotides which comprise the sequences according to  
20 the invention are suitable as hybridization probes for RNA, cDNA and DNA, in order to isolate, in the full length, nucleic acids, or polynucleotides or genes which code for the transcription activator MetR and/or O-succinylhomoserine sulfhydrylase or to isolate those  
25 nucleic acids or polynucleotides or genes which have a high similarity with the sequence of the transcription activator MetR gene and/or that of the O-succinylhomoserine sulfhydrylase gene.

Polynucleotides which comprise the sequences according to  
30 the invention are furthermore suitable as primers with the aid of which DNA of genes which code for the transcription activator MetR and/or O-succinylhomoserine sulfhydrylase can be prepared by the polymerase chain reaction (PCR).

Such oligonucleotides which serve as probes or primers comprise at least 30, preferably at least 20, very particularly preferably at least 15 successive nucleotides. Oligonucleotides which have a length of at least 40 or 50 nucleotides are also suitable.

"Isolated" means separated out of its natural environment.

"Polynucleotide" in general relates to polyribonucleotides and polydeoxyribonucleotides, it being possible for these to be non-modified RNA or DNA or modified RNA or DNA.

10 The polynucleotides according to the invention include a polynucleotide according to SEQ ID No. 1 or a fragment prepared therefrom and also those which are at least 70%, preferably at least 80% and in particular at least 90% to 95% identical to the polynucleotide according to SEQ ID  
15 No. 1 or a fragment prepared therefrom.

The polypeptides according to the invention include the polypeptides according to SEQ ID No. 2 and SEQ ID No. 3, in particular those with the biological activity of the transcription activator MetR and of O-succinylhomoserine  
20 sulfhydrylase, and also those which are at least 70%, preferably at least 80%, and in particular which are at least 90% to 95% identical to the polypeptides according to SEQ ID No. 2 and SEQ ID No. 3 and have the activities mentioned.

25 "Polypeptides" are understood as meaning peptides or proteins which comprise two or more amino acids bonded via peptide bonds.

The invention moreover provides a process for the fermentative preparation of amino acids, in particular  
30 methionine, using coryneform bacteria which in particular already produce the amino acids, and in which the nucleotide sequences which code for the metR gene and/or

for the metZ gene are attenuated, in particular eliminated or expressed at a low level.

The term "attenuation" in this connection describes the reduction or elimination of the intracellular activity of one or more enzymes (proteins) in a microorganism which are coded by the corresponding DNA, for example by using a weak promoter or using a gene or allele which codes for a corresponding enzyme with a low activity or inactivates the corresponding gene or enzyme (protein), and optionally combining these measures.

The microorganisms which the present invention provides can prepare L-amino acids, in particular methionine, from glucose, sucrose, lactose, fructose, maltose, molasses, starch, cellulose or from glycerol and ethanol. They can be representatives of coryneform bacteria, in particular of the genus *Corynebacterium*. Of the genus *Corynebacterium*, there may be mentioned in particular the species *Corynebacterium glutamicum*, which is known among experts for its ability to produce L-amino acids.

Suitable strains of the genus *Corynebacterium*, in particular of the species *Corynebacterium glutamicum* (*C. glutamicum*), are in particular the known wild-type strains

*Corynebacterium glutamicum* ATCC13032  
*Corynebacterium acetoglutamicum* ATCC15806  
*Corynebacterium acetoacidophilum* ATCC13870  
*Corynebacterium melassecola* ATCC17965  
*Corynebacterium thermoaminogenes* FERM BP-1539  
*Brevibacterium flavum* ATCC14067  
*Brevibacterium lactofermentum* ATCC13869 and  
*Brevibacterium divaricatum* ATCC14020

or L-amino acid-producing mutants or strains prepared therefrom, such as, for example, the L-methionine-producing strain



*Corynebacterium glutamicum* ATCC21608.

The inventors have succeeded in isolating the new metR and metZ genes from *C. glutamicum* which code for the transcription activator MetR and the enzyme O-succinylhomoserine sulfhydrylase.

To isolate the metR gene, the metZ gene or also other genes of *C. glutamicum*, a gene library of this microorganism is first set up in *Escherichia coli* (*E. coli*). The setting up of gene libraries is described in generally known textbooks and handbooks. The textbook by Winnacker: Gene und Klone, Eine Einführung in die Gentechnologie (Verlag Chemie, Weinheim, Germany, 1990), or the handbook by Sambrook et al.: Molecular Cloning, A Laboratory Manual (Cold Spring Harbor Laboratory Press, 1989) may be mentioned as an example. A well-known gene library is that of the *E. coli* K-12 strain W3110 set up in  $\lambda$  vectors by Kohara et al. (Cell 50, 495-508 (1987)). Bathe et al. (Molecular and General Genetics, 252:255-265, 1996) describe a gene library of *C. glutamicum* ATCC13032, which was set up with the aid of the cosmid vector SuperCos I (Wahl et al., 1987, Proceedings of the National Academy of Sciences USA, 84:2160-2164) in the *E. coli* K-12 strain NM554 (Raleigh et al., 1988, Nucleic Acids Research 16:1563-1575).

Börmann et al. (Molecular Microbiology 6(3), 317-326 (1992)) in turn describe a gene library of *C. glutamicum* ATCC13032 using the cosmid pHC79 (Hohn and Collins, Gene 11, 291-298 (1980)). To prepare a gene library of *C. glutamicum* in *E. coli* it is also possible to use plasmids such as pBR322 (Bolivar, Life Sciences, 25, 807-818 (1979)) or pUC9 (Vieira et al., 1982, Gene, 19:259-268). Suitable hosts are, in particular, those *E. coli* strains which are restriction- and recombination-defective. An example of these is the strain DH5 $\alpha$ mcr, which has been described by Grant et al. (Proceedings of the National Academy of Sciences USA, 87 (1990) 4645-4649). The long DNA fragments

cloned with the aid of cosmids can in turn be subcloned in the usual vectors suitable for sequencing and then sequenced, as is described e.g. by Sanger et al.

(Proceedings of the National Academy of Sciences of the  
5 United States of America, 74:5463-5467, 1977).

The resulting DNA sequences can then be investigated with known algorithms or sequence analysis programs, such as e.g. that of Staden (Nucleic Acids Research 14, 217-232 (1986)), that of Marck (Nucleic Acids Research 16, 1829-  
10 1836 (1988)) or the GCG program of Butler (Methods of Biochemical Analysis 39, 74-97 (1998)).

The new DNA sequence of *C. glutamicum* which codes for the metR and metZ genes and which, as SEQ ID No. 1, is a constituent of the present invention has been found in this  
15 manner. The amino acid sequence of the corresponding proteins has furthermore been derived from the present DNA sequence by the methods described above. The resulting amino acid sequences of the metR and metZ gene products are shown in SEQ ID No. 2 and SEQ ID No. 3.

20 Coding DNA sequences which result from SEQ ID No. 1 by the degeneracy of the genetic code are also a constituent of the invention. Conservative amino acid exchanges, such as e.g. exchange of glycine for alanine or of aspartic acid for glutamic acid in proteins, are furthermore known among  
25 experts as "sense mutations" which do not lead to a fundamental change in the activity of the protein, i.e. are of neutral function. It is furthermore known that changes on the N and/or C terminus of a protein cannot substantially impair or can even stabilize the function  
30 thereof. Information in this context can be found by the expert, inter alia, in Ben-Bassat et al. (Journal of Bacteriology 169:751-757 (1987)), in O'Regan et al. (Gene 77:237-251 (1989)), in Sahin-Toth et al. (Protein Sciences 3:240-247 (1994)), in Hochuli et al. (Bio/Technology  
35 6:1321-1325 (1988)) and in known textbooks of genetics and

molecular biology. Amino acid sequences which result in a corresponding manner from SEQ ID No. 2 or SEQ ID No. 3 are also a constituent of the invention.

In the same way, DNA sequences which hybridize with SEQ ID  
5 No. 1 or parts of SEQ ID No. 1 are a constituent of the invention. Finally, DNA sequences which are prepared by the polymerase chain reaction (PCR) using primers which result from SEQ ID No. 1 are a constituent of the invention. Such oligonucleotides typically have a length  
10 of at least 15 nucleotides.

Instructions for identifying DNA sequences by means of hybridization can be found by the expert, inter alia, in the handbook "The DIG System Users Guide for Filter Hybridization" from Boehringer Mannheim GmbH (Mannheim,  
15 Germany, 1993) and in Liebl et al. (International Journal of Systematic Bacteriology (1991) 41: 255-260). The hybridization takes place under stringent conditions, i.e. only hybrids in which the probe and target sequence, i.e. the polynucleotides treated with the probe, are at least  
20 70% identical are formed. It is known that the stringency of the hybridization, including the washing steps, is influenced or determined by varying the buffer composition, the temperature and the salt concentration. The hybridization reaction is preferably carried out under a  
25 relatively low stringency compared with the washing steps (Hybaid Hybridisation Guide, Hybaid Limited, Teddington, UK, 1996).

A 5x SSC buffer at a temperature of approx. 50 - 68°C, for example, can be employed for the hybridization reaction.  
30 Probes can also hybridize here with polynucleotides which are less than 70% identical to the sequence of the probe. Such hybrids are less stable and are removed by washing under stringent conditions. This can be achieved, for example, by lowering the salt concentration to 2x SSC and  
35 optionally subsequently 0.5x SSC (The DIG System User's

Guide for Filter Hybridisation, Boehringer Mannheim, Mannheim, Germany, 1995) a temperature of approx. 50 - 68°C being established. It is optionally possible to lower the salt concentration to 0.1x SSC. Polynucleotide fragments which are, for example, at least 70% or at least 80% or at least 90% to 95% identical to the sequence of the probe employed can be isolated by increasing the hybridization temperature stepwise from 50 to 68°C in steps of approx. 1 - 2°C. Further instructions on hybridization are obtainable on the market in the form of so-called kits (e.g. DIG Easy Hyb from Roche Diagnostics GmbH, Mannheim, Germany, Catalogue No. 1603558).

Instructions for amplification of DNA sequences with the aid of the polymerase chain reaction (PCR) can be found by the expert, inter alia, in the handbook by Gait: Oligonucleotide synthesis: A Practical Approach (IRL Press, Oxford, UK, 1984) and in Newton and Graham: PCR (Spektrum Akademischer Verlag, Heidelberg, Germany, 1994).

During work on the present invention, it was found that coryneform bacteria produce amino acids, in particular L-methionine, in an improved manner after attenuation of the metR and/or metZ gene.

To achieve an attenuation, either the expression of the metR and/or of the metZ gene or the catalytic properties of the enzyme proteins can be reduced or eliminated. The two measures can optionally be combined.

The reduction in gene expression can take place by suitable culturing or by genetic modification (mutation) of the signal structures of gene expression. Signal structures of gene expression are, for example, repressor genes, activator genes, operators, promoters, attenuators, ribosome binding sites, the start codon and terminators. The expert can find information on this e.g. in the patent application WO 96/15246, in Boyd and Murphy (Journal of Bacteriology 170: 5949 (1988)), in Voskuil and Chambliss

- (Nucleic Acids Research 26: 3548 (1998), in Jensen and Hammer (Biotechnology and Bioengineering 58: 191 (1998)), in Pátek et al. (Microbiology 142: 1297 (1996)), Vasicova et al. (Journal of Bacteriology 181: 6188 (1999)) and in
- 5 known textbooks of genetics and molecular biology, such as e.g. the textbook by Knippers ("Molekulare Genetik", 6th edition, Georg Thieme Verlag, Stuttgart, Germany, 1995) or that by Winnacker ("Gene und Klone", VCH Verlagsgesellschaft, Weinheim, Germany, 1990).
- 10 Mutations which lead to a change or reduction in the catalytic properties of enzyme proteins are known from the prior art; examples which may be mentioned are the works by Qiu and Goodman (Journal of Biological Chemistry 272: 8611-8617 (1997)), Sugimoto et al. (Bioscience Biotechnology and
- 15 Biochemistry 61: 1760-1762 (1997)) and Möckel ("Die Threonindehydratase aus Corynebacterium glutamicum: Aufhebung der allosterischen Regulation und Struktur des Enzyms", Reports from the Jülich Research Centre, Jül-2906, ISSN09442952, Jülich, Germany, 1994). Summarizing
- 20 descriptions can be found in known textbooks of genetics and molecular biology, such as e.g. that by Hagemann ("Allgemeine Genetik", Gustav Fischer Verlag, Stuttgart, 1986).
- Possible mutations are transitions, transversions,
- 25 insertions and deletions. Depending on the effect of the amino acid exchange on the enzyme activity, "missense mutations" or "nonsense mutations" are referred to. Insertions or deletions of at least one base pair in a gene lead to "frame shift mutations", as a consequence of which
- 30 incorrect amino acids are incorporated or translation is interrupted prematurely. Deletions of several codons typically lead to a complete loss of the enzyme activity. Instructions on generation of such mutations are prior art and can be found in known textbooks of genetics and
- 35 molecular biology, such as e.g. the textbook by Knippers ("Molekulare Genetik", 7th edition, Georg Thieme Verlag,

Stuttgart, Germany, 1997), that by Winnacker ("Gene und Klone", VCH Verlagsgesellschaft, Weinheim, Germany, 1999) or that by Hagemann ("Allgemeine Genetik", 4th edition, Gustav Fischer Verlag, Stuttgart, 1999).

- 5 A common method of mutating genes of *C. glutamicum* is the method of "gene disruption" and "gene replacement" described by Schwarzer and Pühler (Bio/Technology 9, 84-87 (1991)).

In the method of gene disruption a central part of the  
10 coding region of the gene of interest is cloned in a plasmid vector which can replicate in a host (typically *E. coli*), but not in *C. glutamicum*. Possible vectors are, for example, pSUP301 (Simon et al., Bio/Technology 1, 784-791 (1983)), pK18mob or pK19mob (Schäfer et al., Gene 145, 69-  
15 73 (1994)), pK18mobsacB or pK19mobsacB (Jäger et al., Journal of Bacteriology 174: 5462-65 (1992)), pGEM-T (Promega corporation, Madison, WI, USA), pCR2.1-TOPO (Shuman (1994). Journal of Biological Chemistry 269:32678-84; US Patent 5,487,993), pCR®Blunt (Invitrogen, Groningen,  
20 The Netherlands; Bernard et al., Journal of Molecular Biology, 234: 534-541 (1993)) or pEM1 (Schrumpf et al., 1991, Journal of Bacteriology 173:4510-4516). The plasmid vector which contains the central part of the coding region of the gene is then transferred into the desired strain of  
25 *C. glutamicum* by conjugation or transformation. The method of conjugation is described, for example, by Schäfer et al. (Applied and Environmental Microbiology 60, 756-759 (1994)). Methods for transformation are described, for example, by Thierbach et al. (Applied Microbiology and  
30 Biotechnology 29, 356-362 (1988)), Dunican and Shivnan (Bio/Technology 7, 1067-1070 (1989)) and Tauch et al. (FEMS Microbiological Letters 123, 343-347 (1994)). After homologous recombination by means of a "cross over" event, the coding region of the gene in question is interrupted by  
35 the vector sequence and two incomplete alleles are obtained, one lacking the 3' end and one lacking the 5'

end. This method has been used, for example, by Fitzpatrick et al. (Applied Microbiology and Biotechnology 42, 575-580 (1994)) to eliminate the recA gene of *C. glutamicum*.

5 In the method of "gene replacement", a mutation, such as e.g. a deletion, insertion or a base exchange, is established in vitro in the gene of interest. The allele prepared is in turn cloned in a vector which is not replicative for *C. glutamicum* and this is then transferred  
10 into the desired host of *C. glutamicum* by transformation or conjugation. After homologous recombination by means of a first "cross over" event which effects integration and a suitable second "cross-over" event which effects excision in the target gene or in the target sequence, the  
15 incorporation of the mutation or of the allele is achieved. This method was used, for example, by Peters-Wendisch et al. (Microbiology 144, 915 - 927 (1998)) to eliminate the pyc gene of *C. glutamicum* by a deletion.

A deletion, insertion or a base exchange can be  
20 incorporated into the metR gene or the metZ gene in this manner.

In addition, it may be advantageous for the production of L-amino acids, in particular L-methionine, to enhance, in particular to over-express, one or more enzymes of the  
25 particular biosynthesis pathway, of glycolysis, of anaplerosis, of the citric acid cycle or of amino acid export, in addition to attenuation of the metR gene and/or of the metZ gene.

Thus, for example, for the preparation of L-methionine, one  
30 or more genes chosen from the group consisting of

- the gap gene which codes for glyceraldehyde 3-phosphate dehydrogenase (Eikmanns (1992), Journal of Bacteriology 174:6076-6086),

- the tpi gene which codes for triose phosphate isomerase (Eikmanns (1992), Journal of Bacteriology 174:6076-6086),
- the pgk gene which codes for 3-phosphoglycerate kinase (Eikmanns (1992), Journal of Bacteriology 174:6076-6086),
- 5 • the pyc gene which codes for pyruvate carboxylase (Peters-Wendisch et al. (Microbiology 144, 915 - 927 (1998)),
- the lysC gene which codes for a feed-back resistant aspartate kinase (Accession No.P26512),
- 10 • the metA gene which codes for homoserine O-acetyltransferase (ACCESSION Number AF052652),
- the metB gene which codes for cystathionine gamma-synthase (ACCESSION Number AF126953),
- the aecD gene which codes for cystathionine gamma-lyase
- 15 (ACCESSION Number M89931),
- the glyA gene which codes for serine hydroxymethyltransferase (JP-A-08107788),
- the metY gene which codes for O-acetylhomoserine sulfhydrylase (DSM 13556)
- 20 can be enhanced, in particular over-expressed.

It may furthermore be advantageous for the production of L-methionine, in addition to the attenuation of the metR and/or metZ gene, for one or more genes chosen from the group consisting of

- 25 • the thrB gene which codes for homoserine kinase (ACCESSION Number P08210),
- the ilvA gene which codes for threonine dehydratase (ACCESSION Number Q04513),



- the thrC gene which codes for threonine synthase (ACCESSION Number P23669),
- the ddh gene which codes for meso-diaminopimelate D-dehydrogenase (ACCESSION Number Y00151)
- 5 • the pck gene which codes for phosphoenol pyruvate carboxykinase (DE 199 50 409.1; DSM 13047),
- the pgi gene which codes for glucose 6-phosphate isomerase (US 09/396,478; DSM 12969),
- the poxB gene which codes for pyruvate oxidase (DE: 1995  
10 1975.7)

to be attenuated, in particular for the expression thereof to be reduced.

In addition to attenuation of the metR gene and/or of the metZ gene it may furthermore be advantageous, for the  
15 production of amino acids, in particular L-methionine, to eliminate undesirable side reactions (Nakayama: "Breeding of Amino Acid Producing Micro-organisms", in: Overproduction of Microbial Products, Krumphanzl, Sikyta, Vanek (eds.), Academic Press, London, UK, 1982).

20 The invention also provides the microorganisms prepared according to the invention, and these can be cultured continuously or discontinuously in the batch process (batch culture) or in the fed batch (feed process) or repeated fed batch process (repetitive feed process) for the purpose of  
25 production of L-amino acids, in particular L-methionine. A summary of known culture methods are described in the textbook by Chmiel (Bioprozesstechnik 1. Einführung in die Bioverfahrenstechnik (Gustav Fischer Verlag, Stuttgart, 1991)) or in the textbook by Storhas (Bioreaktoren und  
30 periphere Einrichtungen (Vieweg Verlag, Braunschweig/Wiesbaden, 1994)).

The culture medium to be used must meet the requirements of the particular strains in a suitable manner. Descriptions of culture media for various microorganisms are contained in the handbook "Manual of Methods for General  
5 Bacteriology" of the American Society for Bacteriology (Washington D.C., USA, 1981).

Sugars and carbohydrates, such as e.g. glucose, sucrose, lactose, fructose, maltose, molasses, starch and cellulose, oils and fats, such as, for example, soya oil, sunflower  
10 oil, groundnut oil and coconut fat, fatty acids, such as, for example, palmitic acid, stearic acid and linoleic acid, alcohols, such as, for example, glycerol and ethanol, and organic acids, such as, for example, acetic acid, can be used as the source of carbon. These substances can be used  
15 individually or as a mixture.

Organic nitrogen-containing compounds, such as peptones, yeast extract, meat extract, malt extract, corn steep liquor, soya bean flour and urea, or inorganic compounds, such as ammonium sulfate, ammonium chloride, ammonium  
20 phosphate, ammonium carbonate and ammonium nitrate, can be used as the source of nitrogen. The sources of nitrogen can be used individually or as a mixture.

Phosphoric acid, potassium dihydrogen phosphate or dipotassium hydrogen phosphate or the corresponding sodium-  
25 containing salts can be used as the source of phosphorus. The culture medium must furthermore comprise salts of metals, such as, for example, magnesium sulfate or iron sulfate, which are necessary for growth. Finally, essential growth substances, such as amino acids and  
30 vitamins, can be employed in addition to the abovementioned substances. Suitable precursors can moreover be added to the culture medium. The starting substances mentioned can be added to the culture in the form of a single batch, or can be fed in during the culture in a suitable manner.

Basic compounds, such as sodium hydroxide, potassium hydroxide, ammonia or aqueous ammonia, or acid compounds, such as phosphoric acid or sulfuric acid, can be employed in a suitable manner to control the pH of the culture.

5 Antifoams, such as, for example, fatty acid polyglycol esters, can be employed to control the development of foam. Suitable substances having a selective action, such as, for example, antibiotics, can be added to the medium to maintain the stability of plasmids. To maintain aerobic  
10 conditions, oxygen or oxygen-containing gas mixtures, such as, for example, air, are introduced into the culture. The temperature of the culture is usually 20°C to 45°C, and preferably 25°C to 40°C. Culturing is continued until a maximum of the desired product has formed. This target is  
15 usually reached within 10 hours to 160 hours.

Methods for the determination of L-amino acids are known from the prior art. The analysis of L-methionine can be carried out by ion exchange chromatography with subsequent ninhydrin derivatization, as described by Spackman et al.  
20 (Analytical Chemistry, 30, (1958), 1190).

The process according to the invention is used for the fermentative preparation of amino acids, in particular L-methionine.

The present invention is explained in more detail in the  
25 following with the aid of embodiment examples.

The isolation of plasmid DNA from *Escherichia coli* and all techniques of restriction, Klenow and alkaline phosphatase treatment were carried out by the method of Sambrook et al. (Molecular Cloning. A Laboratory Manual (1989) Cold Spring  
30 Harbour Laboratory Press, Cold Spring Harbor, NY, USA). Methods for transformation of *Escherichia coli* are also described in this handbook.

The composition of the usual nutrient media, such as LB or TY medium, can also be found in the handbook by Sambrook et al.

#### Example 1

- 5 Preparation of a genomic cosmid gene library from  
Corynebacterium glutamicum ATCC 13032

Chromosomal DNA from Corynebacterium glutamicum ATCC 13032 was isolated as described by Tauch et al. (1995, Plasmid 33:168-179) and partly cleaved with the restriction enzyme  
10 Sau3AI (Amersham Pharmacia, Freiburg, Germany, Product Description Sau3AI, Code no. 27-0913-02). The DNA fragments were dephosphorylated with shrimp alkaline phosphatase (Roche Diagnostics GmbH, Mannheim, Germany, Product Description SAP, Code no. 1758250). The DNA of the  
15 cosmid vector SuperCos1 (Wahl et al. (1987) Proceedings of the National Academy of Sciences USA 84:2160-2164), obtained from Stratagene (La Jolla, USA, Product Description SuperCos1 Cosmid Vector Kit, Code no. 251301) was cleaved with the restriction enzyme XbaI (Amersham  
20 Pharmacia, Freiburg, Germany, Product Description XbaI, Code no. 27-0948-02) and likewise dephosphorylated with shrimp alkaline phosphatase.

The cosmid DNA was then cleaved with the restriction enzyme BamHI (Amersham Pharmacia, Freiburg, Germany, Product  
25 Description BamHI, Code no. 27-0868-04). The cosmid DNA treated in this manner was mixed with the treated ATCC13032 DNA and the batch was treated with T4 DNA ligase (Amersham Pharmacia, Freiburg, Germany, Product Description T4-DNA-Ligase, Code no.27-0870-04). The ligation mixture was then  
30 packed in phages with the aid of Gigapack II XL Packing Extract (Stratagene, La Jolla, USA, Product Description Gigapack II XL Packing Extract, Code no. 200217).

For infection of the E. coli strain NM554 (Raleigh et al. 1988, Nucleic Acid Research 16:1563-1575) the cells were

taken up in 10 mM MgSO<sub>4</sub> and mixed with an aliquot of the phage suspension. The infection and titering of the cosmid library were carried out as described by Sambrook et al. (1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor), the cells being plated out on LB agar (Lennox, 1955, Virology, 1:190) with 100 mg/l ampicillin. After incubation overnight at 37°C, recombinant individual clones were selected.

### Example 2

#### 10 Isolation and sequencing of the metR and metZ genes

The cosmid DNA of an individual colony was isolated with the Qiaprep Spin Miniprep Kit (Product No. 27106, Qiagen, Hilden, Germany) in accordance with the manufacturer's instructions and partly cleaved with the restriction enzyme  
15 Sau3AI (Amersham Pharmacia, Freiburg, Germany, Product Description Sau3AI, Product No. 27-0913-02). The DNA fragments were dephosphorylated with shrimp alkaline phosphatase (Roche Diagnostics GmbH, Mannheim, Germany, Product Description SAP, Product No. 1758250). After  
20 separation by gel electrophoresis, the cosmid fragments in the size range of 1500 to 2000 bp were isolated with the QiaExII Gel Extraction Kit (Product No. 20021, Qiagen, Hilden, Germany).

The DNA of the sequencing vector pZero-1, obtained from  
25 Invitrogen (Groningen, The Netherlands, Product Description Zero Background Cloning Kit, Product No. K2500-01) was cleaved with the restriction enzyme BamHI (Amersham Pharmacia, Freiburg, Germany, Product Description BamHI, Product No. 27-0868-04). The ligation of the cosmid  
30 fragments in the sequencing vector pZero-1 was carried out as described by Sambrook et al. (1989, Molecular Cloning: A laboratory Manual, Cold Spring Harbor), the DNA mixture being incubated overnight with T4 ligase (Pharmacia Biotech, Freiburg, Germany). This ligation mixture was  
35 then electroporated (Tauch et al. 1994, FEMS Microbiol

Letters, 123:343-7) into the E. coli strain DH5 $\alpha$ MCR (Grant, 1990, Proceedings of the National Academy of Sciences U.S.A., 87:4645-4649) and plated out on LB agar (Lennox, 1955, Virology, 1:190) with 50 mg/l zeocin.

- 5 The plasmid preparation of the recombinant clones was carried out with Biorobot 9600 (Product No. 900200, Qiagen, Hilden, Germany). The sequencing was carried out by the dideoxy chain termination method of Sanger et al. (1977, Proceedings of the National Academy of Sciences U.S.A.,
- 10 74:5463-5467) with modifications according to Zimmermann et al. (1990, Nucleic Acids Research, 18:1067). The "RR dRhodamin Terminator Cycle Sequencing Kit" from PE Applied Biosystems (Product No. 403044, Weiterstadt, Germany) was used. The separation by gel electrophoresis and analysis
- 15 of the sequencing reaction were carried out in a "Rotiphoresis NF Acrylamide/Bisacrylamide" Gel (29:1) (Product No. A124.1, Roth, Karlsruhe, Germany) with the "ABI Prism 377" sequencer from PE Applied Biosystems (Weiterstadt, Germany).
- 20 The raw sequence data obtained were then processed using the Staden program package (1986, Nucleic Acids Research, 14:217-231) version 97-0. The individual sequences of the pZero1 derivatives were assembled to a continuous contig. The computer-assisted coding region analysis was prepared
- 25 with the XNIP program (Staden, 1986, Nucleic Acids Research, 14:217-231).

- The resulting nucleotide sequence is shown in SEQ ID No. 1. Analysis of the nucleotide sequence showed two open reading frames of 567 base pairs and 1146 base pairs, which were
- 30 called the metR gene and metZ gene. The metR gene codes for a protein of 189 amino acids, the metZ gene codes for a protein of 382 amino acids.

## SEQUENCE PROTOCOL

&lt;110&gt; Degussa-Hüls AG

5 &lt;120&gt; New nucleotide sequences which code for the metR and metZ genes

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15 &lt;170&gt; PatentIn Ver. 2.1

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25 &lt;223&gt; metR gene

&lt;220&gt;

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30 &lt;223&gt; metZ gene

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45

## Patent claims

1. An isolated polynucleotide from coryneform bacteria,  
which comprises a polynucleotide sequence which codes  
for the metR and/or metZ genes, chosen from the group  
consisting of
  - a) polynucleotide which is identical to the extent of  
at least 70% to a polynucleotide which codes for a  
polypeptide which comprises the amino acid  
sequence of SEQ ID No. 2,
  - b) polynucleotide which is identical to the extent of  
at least 70% to a polynucleotide which codes for a  
polypeptide which comprises the amino acid  
sequence of SEQ ID No. 3,
  - c) polynucleotide which codes for a polypeptide which  
comprises an amino acid sequence which is  
identical to the extent of at least 70% to the  
amino acid sequence of SEQ ID No. 2,
  - d) polynucleotide which codes for a polypeptide which  
comprises an amino acid sequence which is  
identical to the extent of at least 70% to the  
amino acid sequence of SEQ ID No. 3,
  - e) polynucleotide which is complementary to the  
polynucleotides of a), b), c) or d), and
  - f) polynucleotide comprising at least 15 successive  
nucleotides of the polynucleotide sequence of a),  
b), c), d) or e).
2. A polynucleotide as claimed in claim 1, wherein the  
polynucleotide is a preferably recombinant DNA which  
is capable of replication in coryneform bacteria.
3. A polynucleotide as claimed in claim 1, wherein the  
polynucleotide is an RNA.

4. A polynucleotide as claimed in claim 2, comprising the nucleic acid sequence as shown in SEQ ID No. 1.
5. A DNA as claimed in claim 2 which is capable of replication, comprising
  - 5 (i) the nucleotide sequence shown in SEQ ID No. 1, or
  - (ii) at least one sequence which corresponds to sequence (i) within the range of the degeneration of the genetic code, or
  - (iii) at least one sequence which hybridizes with the  
10 sequence complementary to sequence (i) or (ii), and optionally
  - (iv) sense mutations of neutral function in (i).
6. A DNA as claimed in claim 5 which is capable of replication, wherein the hybridization of sequence  
15 (iii) is carried out under a stringency corresponding to at most 2x SSC.
7. A polynucleotide sequence as claimed in claim 1, which codes for a polypeptide which comprises the amino acid sequence in SEQ ID No. 2 and/or SEQ ID No. 3.
- 20 8. A coryneform bacterium in which the metR gene and/or metZ gene is or are attenuated, in particular eliminated.
9. A process for the fermentative preparation of L-amino acids, in particular L-methionine, which comprises  
25 carrying out the following steps:
  - a) fermentation of the coryneform bacteria which produce the desired L-amino acid and in which at least the metR and/or metZ gene or nucleotide sequences which code for them are attenuated;

- b) concentration of the L-amino acid in the medium or in the cells of the bacteria, and
  - c) isolation of the L-amino acid.
10. A process as claimed in claim 9, wherein bacteria in which further genes of the biosynthesis pathway of the desired L-amino acid are additionally enhanced are employed.
11. A process as claimed in claim 9, wherein bacteria in which the metabolic pathways which reduce the formation of the desired amino acid are at least partly eliminated are employed.
12. A process as claimed in claim 9, wherein the expression of the polynucleotide(s) which code(s) for the metR and/or the metZ gene is reduced or attenuated.
13. A process as claimed in claim 9, wherein the catalytic properties of the polypeptides (enzyme protein) for which the polynucleotides metR and/or metZ code are attenuated.
14. A process as claimed in claim 9, wherein for the preparation of L-amino acids, in particular L-methionine, coryneform microorganisms in which at the same time one or more of the genes chosen from the group consisting of
- 14.1 the lysC gene which codes for a feed back resistant aspartate kinase,
  - 14.2 the gap gene which codes for glycerolaldehyde 3-phosphate dehydrogenase,
  - 14.3 the pyc' gene which codes for pyruvate carboxylase,

- 14.4 the tpi gene which codes for triose phosphate isomerase
- 14.5 the metA gene which codes for homoserine O-acetyltransferase
- 5 14.6 the metB gene which codes for cystathionine gamma-synthase
- 14.7 the pgk gene which codes for 3-phosphoglycerate kinase
- 14.8 the aecD gene which codes for cystathionine gamma-lyase
- 10 14.9 the glyA gene which codes for serine hydroxymethyltransferase
- 14.10 the metY gene which codes for O-acetylhomoserine sulfhydrylase
- 15 is or are enhanced or over-expressed are fermented.
- 15. A process as claimed in claim 9, wherein for the preparation of L-amino acids, in particular L-methionine, coryneform microorganisms in which at the same time one or more of the genes chosen from the group consisting of
- 20 15.1 the thrB gene which codes for homoserine kinase
- 15.2 the ilvA gene which codes for threonine dehydratase
- 25 15.3 the thrC gene which codes for threonine synthase
- 15.4 the ddh gene which codes for meso-diaminopimelate D-dehydrogenase



- 15.5 the pck gene which codes for phosphoenol  
pyruvate carboxykinase
- 15.6 the pgi gene which codes for glucose 6-  
phosphate isomerase
- 5 15.7 the poxB gene which codes for pyruvate oxidase  
is or are attenuated are fermented.
16. A coryneform bacterium which contains a vector which  
carries a polynucleotide as claimed in claim 1f.
- 10 17. A process as claimed in one or more of the preceding  
claims, wherein microorganisms of the species  
Corynebacterium glutamicum are employed.
- 15 18. A process for discovering RNA, cDNA and DNA in order  
to isolate nucleic acids, or polynucleotides or genes  
which code for O-succinylhomoserine sulfhydrylase  
(metZ) and/or the transcription activator MetR or have  
a high similarity with the sequence of the O-  
succinylhomoserine sulfhydrylase (metZ) gene or of the  
transcription activator MetR, which comprises  
employing the polynucleotide comprising the  
20 polynucleotide sequences as claimed in claims 1, 2, 3  
or 4 as hybridization probes.

**Abstract**

The invention relates to polynucleotides from coryneform bacteria which code for the metR and/or metZ genes and comprise polynucleotide sequences, chosen from the group  
5 consisting of

- a) polynucleotide which is identical to the extent of at least 70% to a polynucleotide which codes for a polypeptide which comprises the amino acid sequence of SEQ ID No. 2,
- 10 b) polynucleotide which is identical to the extent of at least 70% to a polynucleotide which codes for a polypeptide which comprises the amino acid sequence of SEQ ID No. 3,
- c) polynucleotide which codes for a polypeptide which  
15 comprises an amino acid sequence which is identical to the extent of at least 70% to the amino acid sequence of SEQ ID No. 2,
- d) polynucleotide which codes for a polypeptide which comprises an amino acid sequence which is  
20 identical to the extent of at least 70% to the amino acid sequence of SEQ ID No. 3,
- e) polynucleotide which is complementary to the polynucleotides of a), b), c) or d), and
- 25 f) polynucleotide comprising at least 15 successive nucleotides of the polynucleotide sequences of a), b), c), d) or e)

and a process for the fermentative preparation of L-amino acids using coryneform bacteria in which at least the metR gene and/or the metZ gene is present in attenuated form,  
30 and the use of polynucleotides which comprise the sequences according to the invention as hybridization probes.